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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.	
10/599,748	10/06/2006	Joseph R. Garlich	050990.0201.PCUS02	3652	
	POLSINELLI SHALTON FLANIGAN SUELTHAUS PC 700 W. 47TH STREET			EXAMINER	
700 W. 47TH S				RAE, CHARLESWORTH E	
SUITE 1000 KANSAS CITY	E 1000 SAS CITY, MO 64112-1802		ART UNIT	PAPER NUMBER	
	,		1611		
			MAIL DATE	DELIVERY MODE	
			01/10/2008	PAPER	

Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

	Application No.	Applicant(s)			
	10/599,748	GARLICH ET AL.			
Office Action Summary	Examiner	Art Unit			
	Charleswort Rae	1614			
The MAILING DATE of this communication appears on the cover sheet with the correspondence address Period for Reply					
A SHORTENED STATUTORY PERIOD FOR REPLY WHICHEVER IS LONGER, FROM THE MAILING D. Extensions of time may be available under the provisions of 37 CFR 1.1 after SIX (6) MONTHS from the mailing date of this communication. If NO period for reply is specified above, the maximum statutory period of Failure to reply within the set or extended period for reply will, by statute Any reply received by the Office later than three months after the mailing earned patent term adjustment. See 37 CFR 1.704(b).	ATE OF THIS COMMUNICATION 36(a). In no event, however, may a reply be timwill apply and will expire SIX (6) MONTHS from a cause the application to become ABANDONE	N. nely filed the mailing date of this communication. D (35 U.S.C. § 133).			
Status					
1) Responsive to communication(s) filed on 06 O					
,	,—				
3) Since this application is in condition for allowance except for formal matters, prosecution as to the merits is					
closed in accordance with the practice under Ex parte Quayle, 1935 C.D. 11, 453 O.G. 213.					
Disposition of Claims					
4) Claim(s) 1-12 is/are pending in the application 4a) Of the above claim(s) is/are withdray 5) Claim(s) is/are allowed. 6) Claim(s) is/are rejected. 7) Claim(s) is/are objected to. 8) Claim(s) 1-12 are subject to restriction and/or of	wn from consideration.				
Application Papers		• • • • • • • • • • • • • • • • • • •			
9) The specification is objected to by the Examine 10) The drawing(s) filed on is/are: a) acc Applicant may not request that any objection to the Replacement drawing sheet(s) including the correct 11) The oath or declaration is objected to by the Ex	epted or b) objected to by the Eddrawing(s) be held in abeyance. See tion is required if the drawing(s) is obj	e 37 CFR 1.85(a). lected to. See 37 CFR 1.121(d).			
Priority under 35 U.S.C. § 119					
 12) Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f). a) All b) Some * c) None of: 1. Certified copies of the priority documents have been received. 2. Certified copies of the priority documents have been received in Application No. 3. Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)). * See the attached detailed Office action for a list of the certified copies not received. 					
Attachment(s) 1) Notice of References Cited (PTO-892)	4) Interview Summary				
 2) Notice of Draftsperson's Patent Drawing Review (PTO-948) 3) Information Disclosure Statement(s) (PTO/SB/08) Paper No(s)/Mail Date 	Paper No(s)/Mail Da 5) Notice of Informal P 6) Other:				

DETAILED ACTION

Status of the Claims

Claims 1-12 are currently pending in this application and are the subject of the Office action.

Election/Restrictions

Restriction is required under 35 U.S.C. 121 and 372.

This application contains the following inventions or groups of inventions which are not so linked as to form a single general inventive concept under PCT Rule 13.1.

In accordance with 37 CFR 1.499, applicant is required, in reply to this action, to elect a single invention to which the claims must be restricted.

- Group 1. Claim(s) 1-4, 9-10, and 12, drawn to a method of protecting a patient from one or more treatments that trigger apoptosis. If this group is elected, then the below Summarized Species requirement is also required.
- Group II. Claim(s) 5-6, drawn to a method of treating a patient suffering from damage to normal tissue attributable to stress. If this group is elected, then the below Summarized Species requirement is also required.
- Group III. Claims 7 and 11, drawn to a method of sensitizing cancer cells to an inhibitor of the PI3 kinase pathway. If this group is elected, then the below Summarized Species requirement is also required.

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Group IV. Claim 8, drawn to a method of treating apoptosis associated with a medical procedure. If this group is elected, then the below Summarized Species requirement is also required.

The inventions represented above as Groups I-IV relate to a general inventive concept under PCT Rule 13.1 because, under PCT Rule 13.2, they share the same or corresponding technical features. Specifically, the technical feature of Groups I-IV is the composition comprising a PTEN inhibitor. The inventions lack unity, however, as the common technical feature is known in the art (US Patent 6,777,439, especially col. 1, line 29 to col. 9. line 6). Thus, the requirement is proper as the inventions represented above as Groups I-II lack unity of invention under PCT Rule 13.1.

Species Election regarding Groups I-IV

This application contains claims directed to more than one species of the generic inventions. These species are deemed to lack unity of invention because they are not so linked as to form a single general inventive concept under PCT Rule 13.1. For example, the generic inventions encompass a multiplicity of PTEN inhibitor compounds and compositions species, which would reasonably exhibit different/variable pharmacologic and pharmaceutical properties. Also, the inventions encompass a multiplicity of cancer cells, cancer diseases, and a multiplicity of treatments that trigger apoptosis. The therapeutic effects to be achieved with these different PTEN inhibitor compounds/compositions would reasonably differ substantially depending on the specific compound, as well as the doses and duration of treatment, and the

contemplated targeted end result with respect to the cancer cell species, cancer disease species, or treatments that trigger apoptosis species. Thus, these species are deemed to lack unity of invention because they are not so linked as to form a single general inventive concept under PCT Rule 13.1. In accordance with 37 CFR 1.499, applicant is required, in reply to this action, to elect a single invention to which the claims must be restricted.

Applicant is required to elect a single species from the below list for examination purposes:

- 1) a single disclosed specific chemically defined PTEN inhibitor; and
- 2) a single disclosed disease species.

If applicant elects Group III, then applicant is further required to elect:

3) a single disclosed specific cancer cell type for examination purposes.

If applicant elects Group IV, then applicant is further required to elect:

4) a single disclosed medical procedure for examination purposes.

Applicant is advised that a reply to this requirement <u>must include an identification</u> of the species that is elected consonant with this requirement, and a listing of all claims readable thereon, including any claims subsequently added. An argument that a claim is allowable or that all claims are generic is considered nonresponsive unless accompanied by an election.

Upon the allowance of a generic claim, applicant will be entitled to consideration of claims to the additional species which depend from or otherwise require all the limitations of an allowable generic claim as provided by 37 CFR 1.141. If claims are

added after election, applicant must indicate which are readable upon the elected species (MPEP 809.02(a). Claims 1, 5, 7, 8, 9, and 11 are considered generic to the above species.

Inventorship Notice

Applicant is reminded that upon the cancellation of claims to a non-elected invention, the inventorship must be amended in compliance with 37 CFR 1.48(b) if one or more of the currently named inventors is no longer an inventor of at least one claim remaining in the application. Any amendment of inventorship must be accompanied by a request under 37 CFR 1.48(b) and by the fee required under 37 CFR 1.17(i).

The examiner has required restriction between product and process claims. Where applicant elects claims directed to the product, and the product claims are subsequently found allowable, withdrawn process claims that depend from or otherwise require all the limitations of the allowable product claim will be considered for rejoinder. All claims directed to a nonelected process invention must require all the limitations of an allowable product claim for that process invention to be rejoined.

In the event of rejoinder, the requirement for restriction between the product claims and the rejoined process claims will be withdrawn, and the rejoined process claims will be fully examined for patentability in accordance with 37 CFR 1.104. Thus, to be allowable, the rejoined claims must meet all criteria for patentability including the requirements of 35 U.S.C. 101, 102, 103 and 112. Until all claims to the elected product are found allowable, an otherwise proper restriction requirement between product claims and process claims may be maintained. Withdrawn process claims that are not

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§ 821.04(b). Additionally, in order to retain the right to rejoinder in accordance with the above policy, applicant is advised that the process claims should be amended during prosecution to require the limitations of the product claims. Failure to do so may result in a loss of the right to rejoinder. Further, note that the prohibition against double patenting rejections of 35 U.S.C. 121 does not apply where the restriction requirement is withdrawn by the examiner before the patent issues. See MPEP § 804.01.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Charlesworth Rae whose telephone number is 571-272-6029. The examiner can normally be reached between 9 a.m. to 5:30 p.m. Monday to Friday.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Ardin Marschel, can be reached at 571-272-0718. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR.

Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http:pair-direct.uspto.gov. Should you have any questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 800-217-9197 (toll-free). If you would like assistance from a USPTO Customer Service Representative or access to the

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automated information system, call 800-786-9199 (IN USA OR CANADA) or 571-272-

1000.

4 January 2007 CER

BRIAN-YONG S. KWON PRIMARY EXAMINER